Reply to Office Action of July 9, 2008

REMARKS

Claims 1-5 and 8-14 are pending. No new matter has been added by way of the present

submission. For instance, claims 6 and 7 have been cancelled. Also, claims 1 and 8-10 have

been amended. Support for the amendment of claim 1 can be found in the original claims 6 and

7. Claims 8 and 9 have been amended for clarity. Support for the amendment of claim 10 can be

found in the original claim 7. Thus, no new matter has been introduced into the claims.

In view of the following remarks, Applicants respectfully request that the Examiner

withdraw the rejection under 35 U.S.C. § 103 and allow the currently pending claims.

<u>Issues under 35 U.S.C. § 103(a)</u>

Claims 1-14 stand rejected under 35 U.S.C. § 103(a) as being unpatentable over Kim et

al. (Chem. Eur. J 2001, pp 176), in view of Tajima et al. (Nucleic Acids Res. 2002:2, pp265-

266). Applicants respectfully traverse the rejection.

The Present Invention

The claimed invention is drawn to a preparation for accelerating an exchange reaction

between a nucleotide sequence at a specific site of a double stranded DNA or RNA and its

homologous nucleotide sequence, comprising a cationic polymer of guanidinated poly(L-

lysine)-graft-dextran (guanidinated PLL-g-Dex) having a guanidine group-containing main chain

and a dextran-containing side chain as an active ingredient.

4

MSW/EAC/vas/la

Docket No.: 4600-0130PUS1

Docket No.: 4600-0130PUS1

Distinctions Over the Cited Art

Kim et al. (Chem. Eur. J 2001, pp 176)

The Kim et al. reference teaches the use of a cationic polymer comprised of poly(L-

lysine)graft-dextran. However, Kim et al. do not teach a poly(L-lysine)graft-dextran polymer

having a guanidine group on the main chain. Kim et al. do not teach or suggest the addition of a

guanidine group to said dextran polymer for any molecular biology method steps, much less for

accelerating an exchange reaction of a nucleotide sequence. Kim et al. do not provide

predictable guidance for a poly(L-lysine)graft-dextran polymer having a guanidine group on the

main chain and how it would behave under conditions for an exchange reaction between a

nucleotide sequence at a specific site of a double stranded DNA or RNA or properties of said

poly(L-lysine)graft-dextran polymer.

Tajima et al. (Nucleic Acids Res. 2002:2, pp265-266)

In regards to the Tajima et al. reference, the shortcomings of Kim et al. are not

complemented by the disclosure of Tajima et al., since there is no teaching of a guanidine group

on the main chain of a dextran polymer.

The arginine-rich peptides taught by Tajima et al. such as P16, PLR50, PLL16 and

PLL40 are not a comb-type cationic polymer, like those disclosed in Kim et al. or in the present

invention as claimed, but are structurally different polymers, i.e., these are straight-chain

peptides, as shown in Figure 1 (b). Because these references encompass such distinct cationic

polymers, one skilled in the art could not obtain from such a combination any rational reason to

Docket No.: 4600-0130PUSI

arrive at the particular poly(L-lysine)graft-dextran guanidine polymer claimed. Additionally,

none of the references teach the substitutions of known functional groups such as guanidine for

another to obtain predictable results yielding the claimed poly(L-lysine)graft-dextran guanidine

polymer.

The Examiner's attention is also directed to the data in Figure 2 of Tajima et al., wherein

the exchange ratio of the arginine-rich peptides such as P16 and PLR50 are at most several times

higher than the lysine peptides such as PLL40 and PLL16.

In contrast, the guanidinated poly(L-lysine)-graft-dextran (guanidinated PLL-g-Dex) of

the present invention shows a high accelerating effect so as to increase the nucleotide chain-

exchange ratio by several tens to several hundred times over the conventional PLL-g-Dex of the

prior art, such as that taught by Kim et al. (See Figure 8). Neither Kim et al. nor Tajima et al.

provide any information on how a guanidine group behaves when linked to a dextran polymer,

and in particular to a poly(L-lysine)graft-dextran polymer.

Legal Standard for Determining Prima Facie Obviousness

M.P.E.P. § 2143 sets forth the guidelines in determining obviousness. The Examiner has

to take into account the factual inquiries set forth in Graham v. John Deere, 383 U.S. 1, 17, 148

USPQ 459, 467 (1966), which has provided the controlling framework for an obviousness

analysis. The four Graham factors of: determining the scope and content of the prior art;

ascertaining the differences between the prior art and the claims that are at issue; resolving the

6

MSW/EAC/vas/la

level of ordinary skill in the pertinent art; and evaluating any evidence of secondary

considerations (e.g., commercial success; unexpected results). 383 U.S. 1, 17, 148 USPQ 459,

467 (1966). Second, the Examiner has to provide some rationale for determining obviousness,

wherein M.P.E.P. § 2143 set forth some rationales that were set established in the recent decision

of KSR International Co. v Teleflex Inc., 82 USPQ2d 1385 (U.S. 2007).

With respect to the present invention, it is clear that there differences between the prior

art and the newly amended claims. That is, the combination of Kim et al. and Tajima et al. do

not teach or fairly suggest the presently claimed cationic polymer of guanidinated poly(L-

lysine)-graft-dextran (guanidinated PLL-g-Dex) having a guanidine group-containing main chain

and a dextran-containing side chain as an active ingredient. Applicants respectfully submit that

there would be no rational reason for the artisan to modify the combined teachings of Kim et al.

and Tajima et al. to obtain the present invention, and as such, a prima facie case of obviousness

cannot be said to exist.

Furthermore, even assuming arguendo that there exists a prima facie case of obviousness,

Applicants have provided evidence of unexpected results to overcome the prima facie case. As

mentioned above, the guanidinated poly(L-lysine)-graft-dextran (guanidinated PLL-g-Dex) of

the present invention shows a high accelerating effect so as to increase the nucleotide chain-

exchange ratio by several tens to several hundred times over the conventional PLL-g-Dex of the

prior art, such as that taught by Kim et al. (See Figure 8). As such, reconsideration and

7

withdrawal of the rejection are respectfully requested.

MSW/EAC/vas/la

Docket No.: 4600-0130PUS1

CONCLUSION

In view of the above amendment, applicant believes the pending application is in condition for allowance.

Should there be any outstanding matters that need to be resolved in the present application, the Examiner is respectfully requested to contact Eggerton Campbell Reg. No. 51307 at the telephone number of the undersigned below, to conduct an interview in an effort to expedite prosecution in connection with the present application.

If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies to charge payment or credit any overpayment to Deposit Account No. 02-2448 for any additional fees required under 37.C.F.R. §§1.16 or 1.147; particularly, extension of time fees.

Dated: January 9, 2009

Respectfully submitted,

Marc S. Weiner

Registration No.: 32,181

BIRCH, STEWART, KOLASCH & BIRCH, LLP

8110 Gatehouse Road

Suite 100 East

P.O. Box 747

Falls Church, Virginia 22040-0747

(703) 205-8000

Attorney for Applicant